

**510(k) SUBSTANTIAL EQUIVALENCE DETERMINATION  
DECISION SUMMARY  
DEVICE ONLY TEMPLATE**

**A. 510(k) Number:**

K033404

**B. Analyte:**

Urine Toxicology Control Material (Drug Mixture)

**C. Type of Test:**

Control Material

**D. Applicant:**

Bio-Rad Laboratories

**E. Proprietary and Established Names:**

Liquicheck Urine Toxicology Control (Level C1, C2, C3, C4, C2 low opiate and C3 low opiate).

**F. Regulatory Information:**

1. Regulation section:  
21CFR862.3280
2. Classification:  
Class I
3. Product Code:  
DIF
4. Panel:  
91

**G. Intended Use:**

1. Intended use(s):
2. Indication(s) for use:  
Liquicheck Urine Urine Toxicology Controls are intended for use as a quality control urine to monitor the performance of laboratory urine toxicology confirmatory procedures..
3. Special condition for use statement(s):
4. Special instrument Requirements:

**H. Device Description:**

Liquicheck Qualitative Urine Toxicology Controls are prepared from human urine with added drugs of abuse and metabolites of drugs of abuse, preservatives, stabilizers and constituents of animal origin. They are in liquid form. They contain

Amphetamines, barbiturates, benzodiazepines, cannabinoids, cocaine, ethanol, LSD, methadone, methaqualone, opiates, phencyclidine and propoxyphene propoxyphene. The specific drugs within each class are listed in the package insert. Approximate drug levels based on gravimetric and GC analyses are listed in the package insert. The controls are supplied at various levels, (Level C1, C2, C3, C4, C2 low opiate and C3 low opiate).

**I. Substantial Equivalence Information:**

1. Predicate device name(s):

Liquichek Urine Toxicology Control Levels C1, C2, C3, C4

2. Predicate K number(s):

K021384

3. Comparison with predicate:

This control material is similar in composition to the predicate device, except that the new device contains MDMA, MDA and MDEA in levels C2, C3 and C4.

**J. Standard/Guidance Document Referenced (if applicable):**

**K. Test Principle:**

N/A. This 510(k) describes control material only.

**L. Performance Characteristics (if/when applicable):**

1. Analytical performance:

a. *Precision/Reproducibility:*

N/A

b. *Linearity/assay reportable range:*

N/A

c. *Traceability (controls, calibrators, or method):*

Value assignment is based on the average of triplicate analyses of control solutions at 3 reference laboratories using gas chromatography. Approximate GC and gravimetric values are listed in the package insert. The manufacturer recommends that each laboratory using Liquichek controls should use these results only as a reference and establish its own parameters for precision.

Open vial stability at 2-8 ° C is tested at 6 time points, the last of which ( $t_{\text{final}}$ ) extends to 20% longer than the expiration date. Recovery of the sample tested at that time point is compared, by GCMS, to a freshly opened vial. Acceptance criteria are that  $t_{\text{final}}$  must be within +/- 10% of the  $T_{\text{zero}}$  value.

Real-time stability studies are ongoing. Acceptance criteria are defined as recovery values at each time point that are within +/- 10% of the values determined for vials stored at 2-8 or -20 ° C.

